

residues in cattle kidney tissue. The U.S. Department of Agriculture, Food Safety Quality Service (now known as the Food Safety Inspection Service) agreed to report any detectable residues in other edible tissue and to report to FDA only those cattle kidney tissue reports where the streptomycin residue was 2 ppm or more.

Since issuing this CPG, FDA has established tolerances for dihydrostreptomycin (59 FR 41976, August 16, 1994) and streptomycin (58 FR 47210, September 8, 1993). Tolerances are established for residues of dihydrostreptomycin in uncooked, edible tissues of cattle and swine of 2.0 ppm in kidney and 0.5 ppm in other tissues, and 0.125 ppm in milk. (See 21 CFR 556.200.) Tolerances are established for residues of streptomycin in uncooked, edible tissues of chickens, swine, and calves of 2.0 ppm in kidney, and 0.5 ppm in other tissues. (See 21 CFR 556.610.)

FDA is withdrawing CPG 7125.22, in its entirety, to eliminate obsolete compliance policy.

Dated: June 20, 2006.
Margaret O'K. Glavin,
Associate Commissioner for Regulatory Affairs.
 [FR Doc. E6-10671 Filed 7-6-06; 8:45 am]
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Submission for OMB Review; Comment Request

Periodically, the Health Resources and Services Administration (HRSA) publishes abstracts of information collection requests under review by the Office of Management and Budget (OMB), in compliance with the Paperwork Reduction Act of 1995 (44 U.S.C. chapter 35). To request a copy of the clearance requests submitted to OMB for review, call the HRSA Reports Clearance Office on (301) 443-1129.

The following request has been submitted to the Office of Management and Budget for review under the Paperwork Reduction Act of 1995:

Proposed Project: The Health Education Assistance Loan (HEAL) Program: Forms (OMB No. 0915-0034 Extension)

The HEAL program provides federally insured loans to assure the availability of funds for loans to eligible students to pay for their education costs. In order to administer and monitor the HEAL program, the following forms are utilized: The Application for Contract of Federal Loan Insurance form (used by lenders to make application to the HEAL insurance program and formerly entitled Lenders Application for Contract of Federal Loan Insurance form); the Borrower's Deferment Request form (used by borrowers to request deferments on HEAL loans and used by lenders to determine borrower's eligibility for deferment); the Borrower Loan Status update electronic submission (submitted monthly by lenders to the Secretary on the status of each loan); and the Loan Purchase/Consolidation electronic submission (submitted by lenders to the Secretary to report sales, and purchases of HEAL loans).

The estimates of burden for the forms are as follows:

Collection activity	Number of respondents	Responses per respondent	Total responses	Hours per response	Total burden hours
Application for Contract of Federal Loan Insurance	17	1	17	8 min	3
Borrower's Deferment Request:					
Borrowers	436	1	436	10 min	73
Employers	261	1.669	436	5 min	36
Borrower Loan Status Update	8	18	144	10 min	24
Loan Purchase/Consolidation	17	248	4,216	4 min	281
Total	739	5,249	417

Written comments and recommendations concerning the proposed information collection should be sent within 30 days of this notice to: John Kraemer, Human Resources and Housing Branch, Office of Management and Budget, New Executive Office Building, Room 10235, Washington, DC 20503.

Dated: June 29, 2006.

Cheryl R. Dammons,
Director, Division of Policy Review and Coordination.

[FR Doc. E6-10591 Filed 7-6-06; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Office of the Director, National Institutes of Health; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), notice is hereby given of a meeting of the Advisory Committee to the Director, National Institutes of Health (NIH).

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material,

and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal property.

Name of Committee: Advisory Committee to the Director, NIH.

Date: August 17, 2006.

Time: 2 p.m. to 3 p.m.

Place: National Institutes of Health, Building 31, 31 Center Drive, Room 5B64, Bethesda, MD 20892.

Agenda: To review and evaluate grant applications (Telephone Conference Call).

Contact Person: Shelly Pollard, ACD Coordinator, National Institutes of Health, 9000 Rockville Pike, Building 31, Room 5B64, Bethesda, MD 20892, (301) 496-0959, pollards@mail.nih.gov.